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| APPLICATION NO. | FI   | LING DATE  | FIRST NAMED INVENTOR | ATTORNEY DOCKET NO. | CONFIRMATION NO. |
|-----------------|--|------------|----------------------|---------------------|------------------|
| 09/944,091      | 08/31/2001   |            | David R. Kranz       | 12942.0067.N        | 1349             |
| 26361           | 7590   | 02/12/2004 |                      | EXAMINER            |                  |
| STEPHEN         |  |            | STRICKLAND, JONAS N  |                     |                  |
|                 | HOWREY, SIMON, ARNOLD & WHITE, LLP<br>750 BERING DRIVE |            |                      |                     | PAPER NUMBER     |
| HOUSTON,        | TX 770   | 57         | 1754                 |                     |                  |

DATE MAILED: 02/12/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

| a see ea  | Application No.   | Applicant(s)   |  |  |  |  |  |
|---|---|--|--|--|--|--|--|
|   | 09/944,091  | KRANZ, DAVID R.  |  |  |  |  |  |
| Office Action Summary   | Examiner  | Art Unit   |  |  |  |  |  |
| ·   | Jonas N. Strickland   | 1754   |  |  |  |  |  |
| The MAILING DATE of this communication app<br>Period for Reply  | ears on the cover sheet with the c  | orrespondence address  |  |  |  |  |  |
| A SHORTENED STATUTORY PERIOD FOR REPLY THE MAILING DATE OF THIS COMMUNICATION.  - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication.  - If the period for reply specified above is less than thirty (30) days, a reply - If NO period for reply is specified above, the maximum statutory period w - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b). | 36(a). In no event, however, may a reply be time within the statutory minimum of thirty (30) day will apply and will expire SIX (6) MONTHS from cause the application to become ABANDONE                          | nely filed s will be considered timely. the mailing date of this communication. D (35 U.S.C. § 133). |  |  |  |  |  |
| Status  |   |  |  |  |  |  |  |
| 1)⊠ Responsive to communication(s) filed on the re  | esponse filed on 9/15/03.   | *  |  |  |  |  |  |
| 2a) This action is <b>FINAL</b> . 2b) ⊠ This  | This action is <b>FINAL</b> . 2b)⊠ This action is non-final.  |  |  |  |  |  |  |
|   | Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213. |  |  |  |  |  |  |
| Disposition of Claims   |   |  |  |  |  |  |  |
| 4) ⊠ Claim(s) 1-21 is/are pending in the application. 4a) Of the above claim(s) is/are withdray 5) □ Claim(s) is/are allowed. 6) ⊠ Claim(s) 1-21 is/are rejected. 7) □ Claim(s) is/are objected to. 8) □ Claim(s) are subject to restriction and/or   | vn from consideration.  |  |  |  |  |  |  |
| Application Papers  |   |  |  |  |  |  |  |
| 9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) acce Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct   | epted or b) objected to by the drawing(s) be held in abeyance. See  | e 37 CFR 1.85(a).  |  |  |  |  |  |
| 11)☐ The oath or declaration is objected to by the Ex   | aminer. Note the attached Office  | Action or form PTO-152.  |  |  |  |  |  |
| Priority under 35 U.S.C. § 119  |   |  |  |  |  |  |  |
| <ul> <li>12) Acknowledgment is made of a claim for foreign</li> <li>a) All b) Some * c) None of:</li> <li>1. Certified copies of the priority documents</li> <li>2. Certified copies of the priority documents</li> <li>3. Copies of the certified copies of the prior application from the International Bureau</li> </ul>   | s have been received.<br>s have been received in Applicati<br>rity documents have been receive  | ion No   |  |  |  |  |  |
| * See the attached detailed Office action for a list  | of the certified copies not receive   | ed.  |  |  |  |  |  |
| Attachment(s)   |   |  |  |  |  |  |  |
| 1) Notice of References Cited (PTO-892)   | 4) Interview Summary  |  |  |  |  |  |  |
| Notice of Draftsperson's Patent Drawing Review (PTO-948)     Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)     Paper No(s)/Mail Date  | Paper No(s)/Mail Do 5) Notice of Informal F 6) Other:   | ate.: Patent Application (PTO-152)   |  |  |  |  |  |

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#### **DETAILED ACTION**

## Response to Amendment

This Detailed Action is in response to the amendment filed on 9/15/2003.
 Claims 1-21 are currently pending.

### Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35
 U.S.C. 102 that form the basis for the rejections under this section made in this
 Office action:

A person shall be entitled to a patent unless -

- (b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.
- 3. Claims 1-7 and 16-19 are rejected under 35 U.S.C. 102(b) as being anticipated by Debbage et al. (US Patent 5,762,885).

Debbage et al discloses an apparatus for removing contaminants from gaseous streams. The reference discloses an apparatus for regenerating a catalyst absorber after contact with a combustion exhaust. The regeneration gas may be comprised of hydrogen and carbon dioxide (col. 5, lines 14-15). However, Debbage et al. also discloses wherein the regeneration gas may also be produced in a regeneration gas generator that reforms methane, a low hydrocarbon to a hydrogen-rich synthesis gas, which also comprises carbon dioxide (col. 6, lines 9-16). Therefore, synthesis gas produced from a gasification unit is used to regenerate a catalyst absorber. The catalyst absorber is comprised of an oxidation catalyst, which is comprised of platinum, palladium, as well as rhodium, which is supported on a high surface area support, such as

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alumina, zirconia, titania, silica, or combinations thereof (col. 3, lines 29-35). The high surface area support may be coated on a ceramic or metal matrix structure (col. 3, lines 49-50). The oxidation catalyst is coated with an absorber, which comprises alkali or alkaline earth mixtures of hydroxides, bicarbonates, and carbonates (col. 3, lines 53-58 and col. 4, lines 7-14). With respect to claim 16, Debbage et al continues to teach a turbine exhaust and wherein a portion of the regeneration synthesis gas is recycled back to the exhaust to produce power in the turbine generator (see Figure 1 and col. 7, lines 40-53). Debbage et al continues to disclose a process using a heat recovery steam generator, with respect to claims 17-19 (col. 4, lines 36-53).

### Claim Rejections - 35 USC § 103

- 4. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:
  - (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 5. Claims 8-15, 20 and 21 are rejected under 35 U.S.C. 103(a) as being unpatentable over Debbage et al. (US Patent 5,762,885) as applied to claims 1-7 and 16-19 above, and further in view of Courty et al. (US Patent 4,088,736).

Applicant claims with respect to claims 8-15, 20 and 21, wherein the synthesis gas is further cleaned in an acid removal unit.

The teachings of Debbage et al. have been discussed with respect to claims 1-7 and 16-19. Debbage et al. teaches a process for treating pollutants

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from a turbine in a power-generating stack. However, Debbage et al. does not teach wherein the synthesis gas is cleaned in an acid gas removal unit.

Courty et al. teaches a process for purifying a gas containing hydrogen sulfide from a gasification unit having carbon dioxide (col. 1, lines 20-25), as well as hydrogen and/or carbon monoxide, synthesis gas (col. 3, lines 60-61). Courty et al. continues to teach wherein the hydrogen sulfide is treated with a mass of zinc oxide (see abstract and col. 1, lines 15-46). Courty et al. continues to teach wherein during regeneration cycles zinc oxide cleans the regeneration gas of gaseous sulfur compounds (col. 4, lines 16-26).

Therefore, it would have been obvious to one of ordinary skill in the art to modify the teachings of Debbage et al., which teaches producing a carbon dioxide and hydrogen regeneration gas from a gasification unit and reducing the amount of pollutants produced from the process, such as hydrogen sulfide, based on the teachings of Courty et al., which teaches a process for reducing hydrogen sulfide from a gasification process by passing the gas comprised of synthesis gas and acidic gases, such as hydrogen sulfide and other sulfur compounds onto a bed of zinc oxide. Such modification would have been obvious to one of ordinary skill in the art, because one of ordinary skill in the art, would have expected a gasification process, which includes reducing pollutants, such as hydrogen sulfide from a gas stream comprised of synthesis gas as taught by Courty et al., to be similarly useful and applicable to a gasification process for wherein synthesis gas is used as a regeneration gas produced from a gasification unit as taught by Debbage et al. While Debbage et al. does not

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teach acid gas removal, Courty clearly teaches wherein acidic gases may be removed from gaseous effluents containing synthesis gases from gasification plants of solid, liquid, and gaseous combustibles (col. 3, lines 57-60).

With respect to claims 10-12, Debbage et al. discloses a shift reactor, a shift catalyst, and wherein the shift catalyst converts carbon monoxide to hydrogen and carbon dioxide (col. 5, lines 28-45).

With respect to claim 13, it would have been obvious to one of ordinary skill in the art to expect the process disclosed by Debbage et al. in view of Courty to convert a carbonyl sulfide to hydrogen sulfide and carbon dioxide, since Debbage et al. teaches a shift catalyst and shift reactor and Courty teaches a gas which comprises COS (col. 3, line 62).

### Response to Arguments

6. Applicant's arguments with respect to claims 1-21 have been considered but are most in view of the new ground(s) of rejection.

Applicant argues that Debbage et al. does not suggest that synthesis gas may be used as a catalyst absorber regeneration stream.

However, Debbage et al. discloses wherein the regeneration gas may also be produced in a regeneration gas generator that reforms methane, a low hydrocarbon to a hydrogen-rich synthesis gas, which also comprises carbon dioxide (col. 6, lines 9-16). Therefore, synthesis gas produced from a gasification unit is used to regenerate a catalyst absorber. The catalyst absorber is comprised of an oxidation-catalyst, which is comprised of platinum, palladium, as well as rhodium, which is supported on a high surface area support, such as

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alumina, zirconia, titania, silica, or combinations thereof (col. 3, lines 29-35). The high surface area support may be coated on a ceramic or metal matrix structure (col. 3, lines 49-50). The oxidation catalyst is coated with an absorber, which comprises alkali or alkaline earth mixtures of hydroxides, bicarbonates, and carbonates (col. 3, lines 53-58 and col. 4, lines 7-14).

#### Conclusion

- 7. The prior art made of record and not relied upon is considered pertinent to applicant's disclosure. US Patent 5,599,758 and US Patent 5,441,990.
- 8. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jonas N. Strickland whose telephone number is 571-272-1359. The examiner can normally be reached on M-TH, 7:30-5:00, off 1st Friday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Stanley Silverman can be reached on 571-272-1358. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

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Information regarding the status of an application may be obtained from

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system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-

free).

Jonas N. Strickland

February 8, 2004